

**CLAIM AMENDMENTS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1-9. (Cancelled).
10. (Currently Amended) A method ~~for the treatment or of treating~~ disorders selected from the group consisting of reduced libido, reduced quality of erections, depression of mood and fatigue, said disorders being caused by andropause by administering to a subject in need thereof diagnosed with andropause and having a disorder selected from the group consisting of reduced libido or sexual drive and reduced quality of erection, depression of mood and fatigue, an effective amount of propionyl L-carnitine in combination with acetyl L-carnitine or one of their pharmaceutically acceptable salts the improvement comprising the lack of increase of blood testosterone levels.
11. (Previously Presented) The method according to claim 10, in which the andropause is caused by aging.
12. (Previously Presented) The method according to claim 10, in which the andropause is caused by chemical or surgical castration.
13. (Cancelled).
14. (Previously Presented) The method according to claim 10, in which the pharmaceutically acceptable salt is selected from the group consisting of: chloride, bromide, orotate, acid aspartate, acid citrate, magnesium citrate, acid phosphate, fumarate and acid fumarate, magnesium fumarate, lactate, maleate and acid maleate, mucate, acid oxalate, pamoate, acid pamoate, acid sulphate, glucose phosphate, glucose phosphate, tartrate, acid tartrate, magnesium tartrate, 2-aminoethane sulphonate, magnesium 2-aminoethane sulphonate, choline

tartrate and trichloroacetate.

15. (Previously Presented) The method according to claim 10, in which propionyl L-carnitine in combination with acetyl L-carnitine are in a unit dosage form containing from 4.0 g to 0.50 g of propionyl L-carnitine inner salt, and from 0.50 g to 4.0 g of acetyl L-carnitine inner salt or an equimolar amount of one of their pharmaceutically acceptable salts.

16. (Previously Presented) The method according to claim 15, in unit dose form containing 2 g of propionyl L-carnitine inner salt and 2 g of acetyl L-carnitine inner salt, or an equimolar amount of one of their pharmaceutically acceptable salts.

17. (Previously Presented) The method according to claim 15, in which propionyl L-carnitine in combination with acetyl L-carnitine are formulated together, as a mixture, or are formulated separately.

18. (Previously Presented) The method of claim 17, in which propionyl L-carnitine in combination with acetyl L-carnitine are in a form suitable for oral or parenteral administration.